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         JAN 16
                 CA/CAplus Company Name Thesaurus enhanced and reloaded
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                 IPC version 2007.01 thesaurus available on STN
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                 CA/CAplus updated with revised CAS roles
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                 CA/CAplus enhanced with patent applications from India
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NEWS 8
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                 PATDPASPC enhanced with Drug Approval numbers
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                 EMBASE enhanced with Clinical Trial Number field
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                 TOXCENTER enhanced with reloaded MEDLINE
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                 CAS Registry Number crossover limit increased from 10,000
                 to 300,000 in multiple databases
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NEWS 19
        MAR 16
                 CASREACT coverage extended
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        MAR 20
                MARPAT now updated daily
NEWS 21
        MAR 22
                LWPI reloaded
                RDISCLOSURE reloaded with enhancements
NEWS 22
        MAR 30
                JICST-EPLUS removed from database clusters and STN
NEWS 23
        APR 02
                GENBANK reloaded and enhanced with Genome Project ID field
NEWS 24 APR'30
NEWS 25 APR 30
                CHEMCATS enhanced with 1.2 million new records
NEWS 26 APR 30
                CA/CAplus enhanced with 1870-1889 U.S. patent records
NEWS 27 APR 30
                INPADOC replaced by INPADOCDB on STN
NEWS 28 MAY 01
                New CAS web site launched
NEWS 29
        MAY 08
                 CA/CAplus Indian patent publication number format defined
NEWS 30 MAY 11
                RDISCLOSURE on STN Easy enhanced with new search and display
                 fields
NEWS EXPRESS
             NOVEMBER 10 CURRENT WINDOWS VERSION IS V8.01c, CURRENT
              MACINTOSH VERSION IS V6.0c(ENG) AND V6.0Jc(JP),
              AND CURRENT DISCOVER FILE IS DATED 25 SEPTEMBER 2006.
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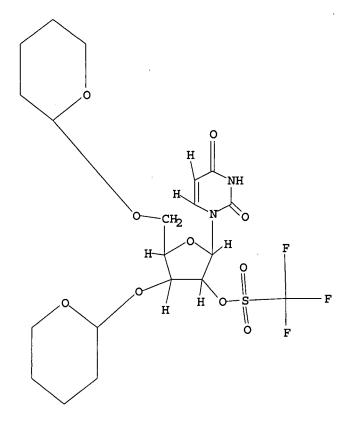
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L1 STRUCTURE UPLOADED

=> d l1

L1 HAS NO ANSWERS

L1 STR



Structure attributes must be viewed using STN Express query preparation.

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=> s 11 sss sam

SAMPLE SEARCH INITIATED 10:42:01 FILE 'REGISTRY'

SAMPLE SCREEN SEARCH COMPLETED - 0 TO ITERATE
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100.0% PROCESSED 0 ITERATIONS

0 ANSWERS

SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE **COMPLETE**

PROJECTED ITERATIONS: BATCH **COMPLETE**

0 TO 0

PROJECTED ANSWERS: 0 TO 0

L2 0 SEA SSS SAM L1

=> s l1 sss full FULL SEARCH INITIATED 10:42:10 FILE 'REGISTRY'

FULL SCREEN SEARCH COMPLETED - 8 TO ITERATE

100.0% PROCESSED 8 ITERATIONS 1 ANSWERS SEARCH TIME: 00.00.01

L3 1 SEA SSS FUL L1

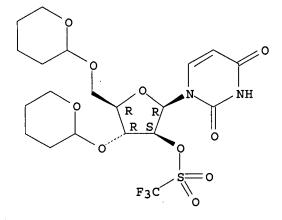
L3 1 ANSWERS REGISTRY COPYRIGHT 2007 ACS on STN

IN 2,4(1H,3H)-Pyrimidinedione, 1-[3,5-bis-O-(tetrahydro-2H-pyran-2-yl)-2-O-[(trifluoromethyl)sulfonyl]- β -D-arabinofuranosyl]- (9CI)

MF C20 H27 F3 N2 O10 S

=> d scan

Absolute stereochemistry.



PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

ALL ANSWERS HAVE BEEN SCANNED

=> file CAplus
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ENTRY SESSION 176.15 176.36

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L1 STRUCTURE UPLOADED

L2 0 S L1 SSS SAM

L3 1 S L1 SSS FULL

FILE 'CAPLUS' ENTERED AT 10:42:28 ON 14 MAY 2007

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=> s 13 ed ibib abs hitstr
MISSING OPERATOR L3 ED
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The search profile that was entered contains terms or nested terms that are not separated by a logical operator.

=> s 13

L41 L3

=> d l4 ed ibib abs hitstr

ANSWER 1 OF 1 CAPLUS COPYRIGHT 2007 ACS on STN L4

ED Entered STN: 22 Oct 2004

ACCESSION NUMBER:

2004:878406 CAPLUS

DOCUMENT NUMBER:

INVENTOR(S):

141:350364

TITLE:

Process for producing 2'-deoxy-2'-fluorouridine Ishi, Akihiro; Ootsuka, Takashi; Kanai, Masatomi; Kuriyama, Yokusu; Yasumoto, Manabu; Inomiya, Kenjin;

Ueda, Koji

PATENT ASSIGNEE(S):

Central Glass Company, Limited, Japan

SOURCE: PCT Int. Appl., 57 pp.

CODEN: PIXXD2

DOCUMENT TYPE:

Patent

LANGUAGE:

Japanese

FAMILY ACC. NUM. COUNT:

PATENT INFORMATION:

PA	PATENT NO.						DATE		APPLICATION NO.				DATE					
WO	VO 2004089968				A1 20041021			WO 2004-JP5109					20040409					
	W:	ΑE,	AG,	AL,	AM,	AT,	AU,	ΑZ,	BA,	BB,	BG,	BR,	BW,	BY,	ΒZ,	CA,	CH,	
		CN,	CO,	CR,	CU,	CZ,	DE,	DK,	DM,	DZ,	EC,	EE,	EG,	ES,	FI,	GB,	GD,	
		GE,	GH,	GM,	HR,	HU,	ID,	IL,	IN,	IS,	KE,	KG,	KΡ,	KR,	KZ,	LC,	LK,	
		LR,	LS,	LT,	LU,	LV,	MA,	MD,	MG,	MK,	MN,	MW,	MX,	MZ,	NA,	NI,	NO,	
		NZ,	OM,	PG,	PH,	PL,	PT,	RO,	RU,	SC,	SD,	SE,	SG,	SK,	SL,	SY,	ТJ,	
		TM,	TN,	TR,	TT,	TZ,	UA,	ŪĠ,	US,	UΖ,	VC,	VN,	YU,	ZA,	ZM,	ZW		
	RW:	BW,	GH,	GM,	ΚE,	LS,	MW,	ΜZ,	SD,	SL,	SZ,	TZ,	ŪĠ,	ZM,	ZW,	AM,	ΑZ,	
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		SK,	TR,	BF,	ВJ,	CF,	CG,	CI,	CM,	GA,	GN,	GQ,	GW,	ML,	MR,	NE,	SN,	
		TD,	TG															
JP	JP 2004323518				A 20041118			JP 2004-115270				20040409						
EP	1612213			A1	20060104			EP 2004-726789				20040409						
•	R:	AT,	BE,	CH,	DE,	DK,	ES,	FR,	GB,	GR,	IT,	LI,	LU,	NL,	SE,	MC,	PT,	
		IE,	SI,	LT,	LV,	FI,	RO,	MK,	CY,	AL,	TR,	BG,	CZ,	EE,	HU,	PL,	SK,	HR
CN	1795	200			Α		2006	0628		CN 2	004-	8001	1679	•	2	0040	409	
US						A1 20061102				US 2005-552463								
PRIORITY APPLN. INFO.:								JP 2003-106849										
									1	WO 2	004-	JP51	9	1	W 2	0040	409	
OTHER S	OTHER SOURCE(S):					CASREACT 141:350364; MARPAT 141:350364												

OTHER S

GI

AB 1-β-D-Arabinofuranosyluracil in a 3',5'-hydroxy-protected form (I; R = hydroxy-protecting group; R1 = H) is reacted with a trifluoromethanesulfonylating agent in the presence of an organic base to convert it into a 2'-triflate form I (R = same as above, R1 = SO2CF3) and this compound is reacted with a fluorinating agent comprising "a salt or complex comprising an organic base and hydrofluoric acid" to produce 2'-deoxy-2'-fluorouridine in a 3',5'-hydroxy-protected form II (R = same as above). An agent for eliminating the protective groups is further caused to act on the protected compound to obtain 2'-deoxy-2'-fluorouridine II (R = H). The 2'-deoxy-2'-fluorouridine obtained can be efficiently purified by temporarily converting it into a 3',5'-diacetyl form, recrystg. the 3',5'-diacetyl form II (R = Ac), and then deacetylating it. Thereby, high-purity 2'-deoxy-2'-fluorouridine can be produced. Thus, 142.90 g I (R = THP, R1 = H), 290 mL DMF, 87.12 g Et3N were added to a SUS pressure vessel, cooled to -54° (inner temperature), treated with 45.00 g CF3SO2F, and warmed to -20° over 2 h 30 min with stirring to give a reaction mixture containing the triflate I (R = THP, R1 = SO2CF3) which was treated with 118.00 g Et3N.3HF at -20° and stirred at room temperature for 62 h 45 min to give, after workup, crude II (R = THP). II (R = THP) (177.18 g) was stirred with 13.80 g p-MeC6H4SO3H.H2O in 150 mL MeOH at room temperature for 16 h 30 min, treated with 6.88 g pyridine, and concentrated to

give crude II (R = H) which was acetylated by 54.10 g Ac20 in 68.46 g pyridine at room temperature for 19 h to give 58.00 g II (R = Ac) (90.17% purity). II (R = Ac) (58.00 g) was recrystd. twice from methanol/H2O (330/120 and 200/100 mL) to give 66% II (R = Ac) (99.95% purity) which (5.00 g) was stirred with 12.89 g NH3 in 50 mL MeOH at room temperature for 6 h 30 min to give 6.77 g high-purity II (R = H).

IT 774611-26-6P

CN

RL: IMF (Industrial manufacture); RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent) (preparation of 2'-deoxy-2'-fluorouridine by triflation of 3',4'-O-protected $\beta\text{-D-arabinofuranosyluracil}$, fluorination, and deprotection)

RN 774611-26-6 CAPLUS

2,4(1H,3H)-Pyrimidinedione, 1-[3,5-bis-O-(tetrahydro-2H-pyran-2-y1)-2-O-[(trifluoromethyl)sulfonyl]- β -D-arabinofuranosyl]- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

REFERENCE COUNT:

THERE ARE 12 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

=> d his

12

L1 L2 L3		FILE 'REGISTRY' ENTERED AT 10:36:34 ON 14 MAY 2007 STRUCTURE UPLOADED 0 S L1 SSS SAM 1 S L1 SSS FULL
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L4		FILE 'CAPLUS' ENTERED AT 10:42:28 ON 14 MAY 2007 1 S L3
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		(FILE 'HOME' ENTERED AT 10:36:21 ON 14 MAY 2007)
L1 L2 L3		FILE 'REGISTRY' ENTERED AT 10:36:34 ON 14 MAY 2007 STRUCTURE UPLOADED 0 S L1 SSS SAM 1 S L1 SSS FULL
L4		FILE 'CAPLUS' ENTERED AT 10:42:28 ON 14 MAY 2007 1 S L3
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